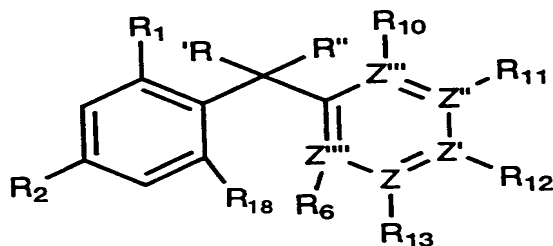
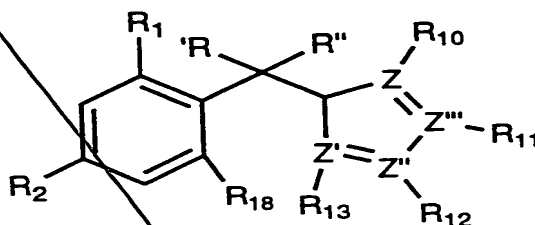


We claim:

1. A compound having the formula:



or



wherein

5  $R_1$  and  $R_2$ , each independently, represent hydrogen or lower alkyl or acyl having 1-4 carbon atoms;

$R'$  and  $R''$  represent hydrogen, lower alkyl or acyl having 1-4 carbon atoms, OH, alkoxy having 1-4 carbon atoms, thiol or thio ether, or amino,

10 or  $R'$  or  $R''$  taken together form an oxo (keto), methano, thioketo, HO-N=, NC-N=,  $(R_7R_8)N-N=$ ,  $R_{17}O-N=$ ,  $R_{17}N=$ , epoxy, cyclopropyl, or cycloalkyl group and wherein the epoxy, cyclopropyl, and cycloalkyl groups can be substituted with lower alkyl having 1-4 carbons or halogen;

15  $R_6$ ,  $R_{10}$ ,  $R_{11}$ ,  $R_{12}$ ,  $R_{13}$  each independently represent hydrogen, a lower alkyl having 1-4 carbons, halogen, nitro,  $OR_7$ ,  $SR_7$ ,  $NR_7R_8$  or  $(CF)_nCF_3$ , and exist only if the Z, Z', Z'', Z''', or Z''' from which it originates is C, or each independently represent hydrogen or a lower alkyl having 1-4 carbons if the Z, Z', Z'', Z''', or Z''' from

which it originates is N, and where one of R<sub>6</sub>, R<sub>10</sub>, R<sub>11</sub>, R<sub>12</sub> or R<sub>13</sub> is X;

R<sub>7</sub> represents hydrogen or a lower alkyl having 1-6 carbons;

R<sub>8</sub> represents hydrogen or a lower alkyl having 1-6 carbons;

5 R<sub>9</sub> represents a lower alkyl having 1-4 carbons, phenyl, aromatic alkyl, or q-hydroxyphenyl, q-bromophenyl, q-chlorophenyl, q-fluorophenyl, or q-iodophenyl, where q=2-4;

R<sub>17</sub> represents hydrogen, lower alkyl having 1-8 carbons, alkenyl (including halogen, acyl, OR<sub>7</sub> and SR<sub>7</sub> substituted alkenes),

10 R<sub>9</sub>, alkyl carboxylic acid (including halogen, acyl, OR<sub>7</sub> and SR<sub>7</sub> substituted alkyls), alkenyl carboxylic acid (including halogen, acyl, OR<sub>7</sub> and SR<sub>7</sub> substituted alkenes), alkyl amines (including halogen, acyl, OR<sub>7</sub> and SR<sub>7</sub> substituted alkyls), and alkenyl amines (including halogen, acyl, OR<sub>7</sub> and SR<sub>7</sub> substituted alkenes);

15 R<sub>18</sub> represents hydrogen, a lower alkyl having 1-4 carbons, halogen, nitro, OR<sub>7</sub>, SR<sub>7</sub>, NR<sub>7</sub>R<sub>8</sub>, or (CF)<sub>n</sub>CF<sub>3</sub>;

X is COOH, tetrazole, PO<sub>3</sub>H, SO<sub>3</sub>H, CHO, CH<sub>2</sub>OH, CONH<sub>2</sub>, COSH, COOR<sub>9</sub>, COSR<sub>9</sub>, CONHR<sub>9</sub>, or COOW where W is a pharmaceutically acceptable salt, and where X can originate from any C or N on the  
20 ring;

Z, Z', Z'', Z''' and Z''', each independently, represent C, S, O, N, or a pharmaceutically acceptable salt, but is not O or S if attached by a double bond to another such Z or if attached to another such Z which is O or S, and is not N if attached by a

25 ~~single bond to another such Z which is N, and~~

*a* n = 0-3,

2. A compound of claim 1 wherein said compound selectively activates Retinoid X Receptors in preference to Retinoic Acid Receptors.

3. A compound selected from the group consisting of 4-  
[1-(2-methyl-4-t-butylphenyl)ethenyl] benzoic acid,  
4-[1-(2-methyl-4-t-butylphenyl)cyclopropyl] benzoic acid,  
4-[(2-methyl-4-t-butylphenyl)carbonyl] benzoic acid,  
5 4-[(2-methyl-4-t-butylphenyl)carbonyl] benzoic acid oxime, and  
4-[1-(2-methyl-4-t-butylphenyl)carbonyl] benzoic acid

methyloxime

4. A pharmaceutical composition comprising in a  
pharmaceutically acceptable vehicle suitable for enteral,  
10 parenteral, or topical administration, one or more compound of  
claim 1.

5. A method for modulating a process mediated by one or  
more Retinoid X Receptors, said method comprising causing said  
process to be conducted in the presence of at least one compound as  
15 set forth in claim 1.

6. A method according to claim 5 wherein said process  
is the *in vivo* modulation of lipid metabolism, *in vivo* modulation of  
skin-related processes, *in vivo* modulation of malignant cell  
development, *in vivo* modulation of premalignant lesions, or *in vivo*  
20 modulation of programmed cell death.

7. A method according to claim 5 wherein said process  
is in *in vivo* or *in vitro* cellular growth and differentiation, or  
*in vivo* limb morphogenesis.

8. A method for modulating a process mediated by one or  
25 more Retinoid X Receptors, said method comprising administering to

a mammalian subject an amount, effective to modulate said process mediated by said one or more Retinoid X Receptors, of one or more compound of claim 1.

5 9. A method for treating a mammalian subject requiring Retinoid X Receptor therapy comprising administering to such subject a pharmaceutically effective amount of one or more compounds as set forth in claim 1.

10 10. A method for increasing plasma concentrations of high density lipoprotein in a mammalian subject comprising administering to such subject a pharmaceutically effective amount of one or more compounds as set forth in claim 1.

15 11. A method for modulating a process mediated by intracellular receptors, said method comprising causing said process to be conducted in the presence of a composition comprising a first compound as set forth in claim 1 which selectively  
20 activates Retinoid X Receptors in preference to Retinoid Acid Receptors, in combination with a second compound which activates one or more intracellular receptors other than Retinoid X Receptors, and wherein the physiological effect in mammals produced by said composition at a given concentration is greater than the additive effect achieved by utilizing each said compound alone at  
~~said concentration~~

add  
E1